TITLE: Preparation of amides and ureas as activators of

soluble guanylate cyclase

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		HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KΖ,	LC,	LK,	LR,	LS,	LT,	
		LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NΖ,	PL,	PT,	RO,	RU,	
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		DE,	DK,	ES,	FΙ,	FR,	GB,	GR,	ΙE,	ΙΤ,	LU,	MC,	NL,	PT,	SE,	TR,	BF,	
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EP	EP 1237849				A1 20020911				EP 2000-973061					20001106				
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OTHER S	HER SOURCE(S).					PAT	134.	3531	75									

OTHER SOURCE(S): MARPAT 134:353175

GΙ

The title compds. R4PZNR1R2 [I; R1, R2 = alkyl; R1R2 together form alkylene; Z = alkylene; P = a direct bond, X, Y, W, XY, YW, XYW (wherein W = O, S, NR3; R3 = H, alkyl; Y = UV; V = a direct bond, alkylene; U = CS, CO, SO2, C(:NR); R = H, OH, alkyl; X = O, NR6; R6 = H, alkyl, alkenyl, etc.); R4 = alkyl, alkenyl, alkynyl, etc.], useful in the activation of soluble guanylate cyclase, were prepared E.g., synthesis of the urea II, starting with 4-bromoaniline and 1-(3-aminopropyl)pyrrolidine, was given. Biol. data for compds. I (e.g., IC50 for inhibition of platelet aggregation) were presented.

IT 338980-58-8P 338980-88-4P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of amides and ureas as activators of soluble guanylate

cyclase)

RN 338980-58-8 ZCAPLUS

CN Benzamide, $2-[[(5-\text{chloro}-1-\text{methyl}-3-\text{phenyl}-1\text{H-pyrazol}-4-\text{yl})\,\text{methyl}]\text{thio}]-N-[3-(dimethylamino)\,\text{propyl}]-(9CI)$ (CA INDEX NAME)

RN 338980-88-4 ZCAPLUS

CN Urea, N-[2-[[(5-chloro-1-methyl-3-phenyl-1H-pyrazol-4-yl)methyl]thio]phenyl]-N'-[3-(dimethylamino)propyl]- (9CI) (CA INDEX NAME)